U.S. Patent Application No. 304 481

## DECLARATIO, N

We, the undersigned Gabrijela Kobrehel and Slobodan Djokić, hereby declare that we have carried out the following comparative tests, whose results are attached to this Declaration:

- Teble 3: Testing of in vitropotency of N-methyl-11-aza10-deoxo-10-dihydroerythromycin A in comparison
  with erythromycin A and its 11-aza derivative against
  182 gram-positive bacterial organisms
- Table 4: Testing of in vitro potency of N-methyl-11-aza-10-deaxo-10-dihydroerythromycin A in comparison with erythromycin A and its 11-aza derivative against 179 gram-negative bacterial organisms
- Table 5: Testing of in vitro sensitivity of 30 anaerobic bacterial organisms against N-methyl-11-aza-10-deoxo-10-dihydroerythromycin A in comparison with erythromycin A and its 11-aza derivative
- Table 6: Testing of acute toxicity in mice by the method of
  Litchfield-Wilcoxon of N-methyl-11-aza-10-deoxo-10dihydroerythromycin A in comparison with erythromycin A
  and its 11-aza derivative
- Table 7: Testing of acid stability at pH 1.2 of N-methyl-11aza-10-deoxo-10-dihydroerythromycin A, 2'-propionyl-N-methyl-11-aza-10-deoxo-10-dihydroerythromycin A,

N-methyl-11-aza-10-deoxo-10-dihydroerythromycin A cyclic 13,14-carbonate, and 2'-acetyl-N-methyl-11-aza-10-deoxo-10-dihydroerythromycin A cyclic 13,14-carbonate in comparison with the parent erythromycin A, by exposure of the compounds to hydrochloric acid (pH 1.2) and determination of the residual activity by two-fold dilution technique vs. Staphilococcus aureus ATCC 6538-P and expressing the results in minimum inhibitory concentrations (MIC) in mcg/ml

Figure 1:Blood level studies in rabbits after oral doses of 100 and 250 mg of N-methyl-11-aza-10-deoxo-10-dihydroerythromycin A (1) in comparison with the oral administration of 250 mg of erythromycin A

The undersigned further declare that all statements made herein of our own knowledge are true and that all statements made on information and belief are believed to be true; and further, that these statements were made with the knowledge that willful, false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful, false statements may jeopardize the validity of the application or any patent issuing thereon.

Zagreb, January 31, 1984

Gabrijela Kolrohel Flaboelem Douic'

Table 3. Antibacterial in vitro activity on gram-negative clinical isolates

TEST					N	IIC (	mcg/i	nl)			
ORGANISM (	COMPOUND	0.5	1.0	2.0	4.0	8.0	16.0	32.0	64.0	128.0 R	of teste strains
Enteroco-	E	26	11	9		6	3		1	7	
ccus	A	20	6	11	ı	5	4		. 4	12	63
	1	15	13	10	6				4	<b>15</b>	
Staph.	E	9	7	12						13	
aureus	A	10	7	12	1			1		10	41
	1	11	10	4	. , 6	3			•	7	
Staph.	E	22	6	12		2					· · · · · · · · · · · · · · · · · · ·
albus	A	24	9	4	2	3	•				42
	1	30	12				:				
Strepto-	E	9	7					· · · · · ·		* <del>***</del> *** *** *** *** ***	
coccus	A	12	4		4				•		16
pneumoniae	1	11	5		···,		•				
Streptoco-	E	17	3								• <del>• • • • • • • • • • • • • • • • • • </del>
coccus	A	20				1.					
haemolyticu	s l	13	. 7								
No. of sensiti	- E	83	× 34	<i>3</i> 4.		8	<del>-</del>		 1	20	182
ve strains	A ·	86	26	26	4	8	4.	1	4	22	
14.4	1	80	47	14	12	3			4	12	

E - Erythromycin A

A - 11-Aza-10-deoxo-10-dihydroerythromycin A

<sup>1 -</sup> N-Methyl-ll-aza-lO-deoxo-lO-dihydroerythromycin A

R - resistant

Table 4. Antibacterial in vitro activity on gram-negative clinical isolates

TEST						M.	IC (mo	g/ml	)		<del></del>
ORGANISM C	OMPOUND	0.5	1.0	2.0	4.0					128.0 R	No.
	<b>E</b> .		1			2	28	40	8	22	100
E. coli	A	l		2	39	50	2	1	2	3	
	1		4	42	45	5	2	1	1	0	
	<u>7</u> .		2	37.	53_	3_		1	2	2_	
	E						1	2	2	4	9
Klebsiella	A.				3	1	3	1		1	. •
pneumon.	ļ				4	4	1				
	7					2	4	1_	2		
•	E			•			1		2	7	10
Klebsiella	A				•	3	5			2	_ =
aerogenes	1					9	•	1		•	
			·			5	1_	3_		1_	
	E			•		13				3	16
Proteus	A			•		2		3	1	(10	~
mirabilis	1				1	2	1	10			
<del>-</del>					2_		<u>l_</u>		66		
r	E							<del></del>	1	9	10
Pseudomo-	A					ı		2	3	4	<b></b> €
nas aerug.	100	× .			٠. ٠	\ <u>.</u> —	~	i i	5.	3	
							1_	4	_1	<u> </u>	
	E		· <b>-</b>		<del></del> ;	7			1	17	18
Enterobac-	A					1	5	7	4	1	<b>20</b> .
ter aerog.	1				3	6	6	2	•	1	
		~~~~~				3_	9	1	_5	·	· · · · · · · · · · · · · · · · · · ·
	 Е						1	<b></b>	- <del></del>	• • • • • • • • • • • • • • • • • • •	1
Enterobac-						1					· : 4
ter liquef					1	_					
<u></u>	7					1_				·	
	E	<b></b>	<del></del>				1	<b></b>		. 2	
Mima	A					• •	_	2	1	. <b>~</b>	3
polymorpha						ı		1	1		
	_					-		_	-		

Continued on the next page:

Table 4 (continued)

TEST						M.	IC (mo	g/ml	)			
ORGANISM (	COMPOUND	0.5	1.0	2.0	4.0	8.0	16.0	32.0	64.0	128.0	R	No.
Herella	E A 1 7		1	1 1	2	1 2	3 1 3	1 3 1	1		1	6
Haemophy-	E A • 1 — 7	1 1 3 1	3 3 4	2 2	3	·						6
No. sensitive strations		1 2 3 1	0 4 8 6	2 4 43 39	3 42 54 57	16 59 28 17	35 16 13 17	45 19 17 11	15 11 7 17		22	179 179 179 179

E - Erythromycin A

A - 11-Aza-10-deoxo-10-dihydroerythromycin A

1 - N-Methyl-ll-aza-10-deoxo-10-dihydroerythromycin A

7 - N-Methyl-ll-aza-10-deoxo-10-dihydroerythromycin A 13,14-cyclic carbonate

Method: Two-fold serial dilution technique in MH agar.

No. - number of tested strains

R - resistant

Table 5. Sensitivity of anaerobic bacteria

TEST			··· · · · · · · · · · · · · · · · · ·		MIC	(mc	g/ml)		··	· · · · · · · · · · · · · · · · · · ·		
ORGANISMS	COMPOUND	0.5	1.0	2.0	4.0	8.0	16.0	32.0	64.0	128.0	R No. of ted	tes-
BACTEROIDES	E			1	3	2						
FRAGILIS	A		1	4	1							6
	1	3		3								
BACTEROIDES	E			2	2	2			4	•		
NECROPHORUS	A	3	1	2	+ 57					•		6
	11	2_	3	1	·		·					
BACTEROIDES	E			2		1	٠					*.*
MELANINOGE-	A	1	1	1		•	•					3
NICUS	1	1_	2_									: ·
VEILONELLA	${f E}$		2	ı	ı							.:
SP	A	3		1					٠			4
	11	<u> 4</u>										·
LEUCONOSTOC	E.		3									
- 	<b>.</b> <b>A</b> .	2	1			_			,			3 🤇
*	11	2	<u> </u>		ij.		<u></u>					
PEPTOSTREPTO-	E	5	3									
coccus	A	7	í									8
	11	8					: ·				<u> </u>	
No.ofsensitive	E	5	8	6	6	5			<b></b>	<del></del>		30
strains	A	16	5	8	1							30
	1	20	6	4	_							30

E - Erythromycin A

Method: Two-fold serial dilution technique in MH agar.

A - 11-Aza-10-deoxo-10-dihydroerythromycin A

<sup>1 -</sup> N-Methyl-ll-aza-10-deoxo-10-dihydroerythromycin A

R - resistant

Table 6. Acute toxicity

Compound+	Route of administration	<sup>LD</sup> 50	<sub>LD</sub> 16	LD <sub>84</sub>
E	I.V. I.P. P.O.	360 (241.9) <sup>++</sup> 520 (349.4)	280 326	360 840 -
A	I.V. I.P. P.O. >	280 (141.7) - 10000 (5060)	220 <u>-</u> .	350 -
1	I.V. I.P. P.O. >	825 (421.6) 1200 (513.2) 10000 (5100)	610 1010	1020

E - Erythromycin A

/cal

A - 11-Aza-10-deoxo-10-dihydroerythromycin A

<sup>1 -</sup> N-Methyl-11-aza-10-deoxo-10-dihydroerythromycin A

The substances were tested as the lactobionate salts.
 The values calculated in the relation to the active compound.

Table 7. Acid stability at pH 1.2 (MIC in mcg/ml)

Exposure time	MIC									
in hours	E	1	5	7	g +					
control-without										
acid	0.1	0.1	0.1	0.5	0.5					
0	5	0.1	0.1	0.5	0.5					
1/2	10	0.1	0.1	0.5	0.5					
1	10	0.5	0.5	0.5	0.5					
3	25	1.0	0.5	1.0	0.5					
6	25	2.5	0.5	0.1	0.5					

E = Erythromycin A

Strain: Staphylococcus aureus ATCC 6538-P

<sup>1 =</sup> N-Methyl-11-aza-10-deoxo-10-dihydroerythromycin A

<sup>5 = 2&#</sup>x27;-Propionyl-N-methyl-ll-aza-10-dexo-10-dihydroerythromycin A

<sup>7 =</sup> N-Methyl-11-aza-10-deoxo-10-dihydroerythromycin A cyclic 13,14-carbonate

<sup>8 = 2&#</sup>x27;-Acetyl-N-methyl-ll-aza-10-dexo-10-dihydroerythromycin A cyclic 13,14-carbonate

<sup>+</sup> Arabic figures correspond to the notation of the Examples

Figure 1. Blood level studies in rabbits after oral doses of 100 and 250 mg

